



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification⁵:

G01N 24/00

A1

(11) International Publication Number:

WO 94/25860

(43) International Publication Date: 10 November 1994 (10.11.94)

(21) International Application Number:

PCT/US94/04822

(22) International Filing Date:

28 April 1994 (28.04.94)

(30) Priority Data:

08/055,050

28 April 1993 (28.04.93)

US

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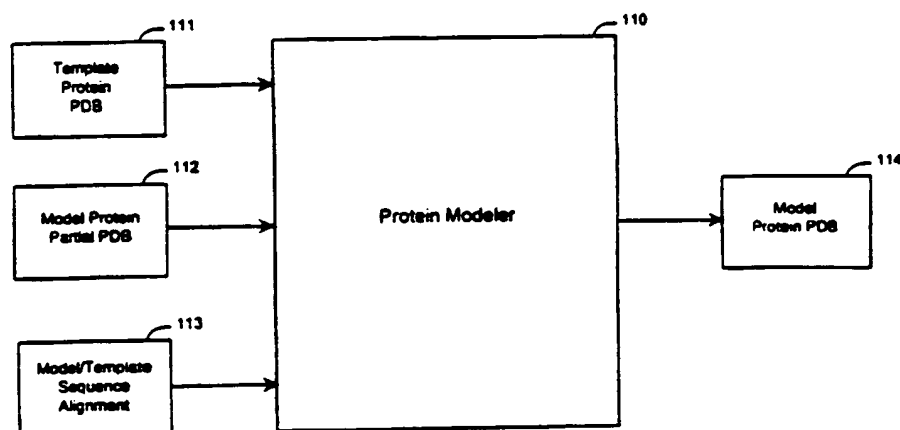
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(81) Designated States: AU, CA, FL, JP, KR, NO, NZ, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).

Published

With international search report.

(54) Title: METHOD AND SYSTEM FOR PROTEIN MODELING



(57) Abstract

A method in a computer system for modeling a three-dimensional structure of a model protein (112) is provided. In a preferred embodiment, the modeling is based upon a three-dimensional structure of a template protein (111) and an amino acid sequence alignment (113) of the model protein and the template protein. The proteins comprise a plurality of amino acids having backbone atoms and side chain atoms. For each amino acid in the model protein, when the template protein has an amino acid aligned with the amino acid of the model protein, the position of each backbone atom of the amino acid of the model protein is established based on the position of a topologically equivalent backbone atom in the aligned amino acid of the template protein. The inter-atomic distance constraints for each pair of atoms with an established position is generated. Finally, the position of each atom in the model protein is set so that the inter-atomic distances are in accordance with the constraints.

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Description

METHOD AND SYSTEM FOR PROTEIN MODELING

Cross-Reference to Prior Application

This application is a continuation-in-part of U.S. Patent Application Serial No. 08/055,050, filed April 28, 1993, now pending.

Technical Field

This invention relates generally to a computer-implemented method and system for modeling the three-dimensional structure of a protein, and, more specifically, to protein modeling using the three-dimensional structure of a template protein and sequence alignment between the protein to be modeled and the template protein.

Background Of The Invention

The function of a protein is related to its three-dimensional structure. The three-dimensional structure of some proteins can be determined using X-ray crystallography or Nuclear Magnetic Resonance (NMR). However, for other proteins, these methods cannot be used to determine the three-dimensional structure. When these methods cannot be used, computer-based protein modeling techniques have been used with some success. These protein modeling techniques use the known three-dimensional structure of a homologous protein to approximate the structure of another protein.

In one such technique, the known three-dimensional structures of the proteins in a given family are superimposed to define the structurally conserved regions in that family. Among the members of a given family, there is considerable variation in the conformations of regions located between two consecutive structurally conserved regions, and thus, these regions are called the variable regions. These variable regions essentially contribute to the identity of a protein in

its family. However, the modeling of the variable regions has been unsatisfactory.

Conventional homology modeling techniques have been used routinely to build models of proteases and antibodies. These techniques generally model the three-dimensional position of amino acids in structurally conserved regions by taking the Cartesian coordinates from homologous amino acids in a template protein with known three-dimensional structure. For the amino acids in the variable regions, these techniques take suitable loops from the Protein Data Bank (PDB). (The PDB is a collection of protein information relating to the known structure of protein. The PDB is administered by Brookhaven National Laboratory.) Although these techniques are generally successful in modeling the structurally conserved regions for structurally undefined members of the family, these techniques have been unsuccessful in modeling of the variable regions. Since variable regions and structurally conserved regions of the model protein come from different protein structures, high energy short contacts are often found in the models. These high energy contacts are usually between inter-variable regions that are grafted from different known protein structures. In cases where the sequence identity in the structurally conserved regions between the template and the model protein is weak, the interior amino acids are also susceptible to short contacts. Generally, these short contacts are removed by performing rotation around single bonds using interactive graphics, which is a tedious, and at times, an impractical procedure. Energy minimization has been used to relax strains in a model. However, the minimization procedure leads to structures that are trapped in local minima and relies entirely on the integrity of the starting structure.

Certain proteins with very weak sequence identities fold into similar three-dimensional structures. For example, the three-dimensional structures of a number of helical cytokines fold in similar three-dimensional topology in spite of weak sequence homology. Members of helical cytokine family

not only show diversity in their disulfide topology but also the disulfide crosslinks are part of the variable regions. Besides the helical cytokines, there are other protein families with weak sequence identities and non-homologous disulfides in the variable regions. The prior homology modeling techniques produce unsatisfactory modeling of these families because of the absence of sequence homology in the structurally conserved regions and the presence of non-homologous disulfide crosslinks in the variable regions.

Summary of the Invention

It is an object of the present invention to provide a method and system for modeling the three-dimensional structure of members of a protein family/superfamily from the known three-dimensional structure of one or more members of that family.

It is another object of the present invention to provide a method and system for modeling the three-dimensional structure of a protein by using position information of all topologically equivalent atoms in a template protein.

It is another object of the present invention to provide a method and system for generating a sequence alignment between two proteins based on the resulting three-dimensional structure model.

It is another object of the present invention to provide a method and system for generating a protein model that minimizes short contacts.

It is another object of the present invention to provide a method and system for modeling proteins with weak sequence identity with a template protein.

It is another object of the present invention to provide a method and system for modeling variable regions between two structurally conserved regions of a protein.

It is another object of the present invention to provide a method and system for modeling variable regions constrained by disulfide crosslinks of a protein.

These and other objects, which will become apparent as the invention is more fully described below, are provided by a computer-implemented method and system for modeling a three-dimensional structure of a model protein. In a preferred embodiment, the modeling is based upon a three-dimensional structure of a template protein and an amino acid sequence alignment of the model protein and the template protein. The proteins comprise a plurality of amino acids having backbone atoms and side chain atoms. For each amino acid in the model protein, when the template protein has an amino acid aligned with the amino acid of the model protein, the position of each backbone atom of the amino acid of the model protein is established based on the position of a topologically equivalent backbone atom in the aligned amino acid of the template protein. Then, the inter-atomic distance constraints for each pair of atoms with an established position is generated. Finally, the position of each atom in the model protein is set so that the inter-atomic distances are in accordance with the constraints.

In another aspect, the present invention provides for a method and system for modeling a three-dimensional structure of a variable region of a model protein. The model protein has amino acids that have positions within a three-dimensional structure. The method receives relative positional information between pairs of amino acids. The relative positional information preferably includes the ψ and ϕ angle values between pairs of amino acids. The method then establishes a position for a first amino acid in the variable region. The method then for each amino acid pair in the variable region generates a model position for the amino acids based on the received relative positional information for the pair of amino acids. In a preferred embodiment the ψ and ϕ angle values are collected for pairs of amino acid combinations in variable regions of a family of proteins. The

generated model position for the amino acids are preferably based on a random selection of the collected ψ and ϕ angle values.

Brief Description Of The Drawings

Figure 1 is an overall block diagram of the input and output for the protein modeler.

Figure 2 is an overall flow diagram of the protein modeler.

Figure 3 is a flow diagram of a routine to generate a TEA position data structure for the model protein.

Figure 4 is a routine to generate distance constraints for topologically equivalent atoms.

Figure 5 is a routine to generate standard constraints for non-topologically atoms.

Figure 6 is a flow diagram showing the DGEOM process.

Figure 7 is a diagram illustrating the ψ and ϕ angles between pairs of amino acids.

Figure 8 is a diagram illustrating the amino acid angle matrix.

Figure 9 is a flow diagram of a function that generates the amino acid angle matrix.

Figure 10 is a flow diagram of a function to generate a variable region model for a model protein based on the amino acid angle matrix.

Detailed Description Of The Invention

The present invention provides a computer-based method and system for modeling the three-dimensional structure of a protein (model protein). In a preferred embodiment, the protein modeler generates the three-dimensional structure of the model protein based on a template protein with a known three-dimensional structure and an amino acid sequence alignment between the model protein and the template protein. Using the template protein and the sequence alignment, the

protein modeler generates a variety of inter-atom distance constraints for conserved regions and standard and chemical constraints for variable regions, and allows for the input of miscellaneous constraints (e.g., disulfide crosslinks). The protein modeler then generates a three-dimensional structure for the model protein using known techniques of distance geometry to ensure compliance with the constraints. Srinivasan and Sudarsanam, Protein Science 2:277-289 (1993), incorporated herein by reference, contains a further description of the present invention.

The protein modeler generates the inter-atomic distance constraints for the model protein based on the position of topologically equivalent atoms (TEA) in the template protein. The protein modeler determines whether each atom in the model protein is topologically equivalent to an atom in the template protein. An atom of an amino acid of the model protein is topologically equivalent when the amino acid of the model protein aligns with an amino acid of the template protein and the amino acid of the template protein has a corresponding atom. If the aligned amino acids are identical, then each atom of the model amino acid corresponds to the atom of the same name in the template amino acid. For example, the C-alpha atom of the model amino acid corresponds to the C-alpha atom of the template amino acid, and the C-beta atom of the model amino acid corresponds to the C-beta atom of the template amino acid. If the aligned amino acids are not identical, then each atom of the model amino acid may or may not have a corresponding atom in the template amino acid. Each backbone atom (N, C-alpha, C', and O) has a corresponding atom. However, the correspondence between side chain atoms depends on the aligned amino acids. For example, if the model amino acid is arginine and the template amino acid is alanine, then the C-beta atoms correspond. However, the C-gamma, C-delta, and other atoms of the arginine side chain have no corresponding atoms in the alanine side chain.

The protein modeler sets the position of each topologically equivalent atom in the model protein to the same position as that of the topologically equivalent atom in the template protein. The protein modeler then calculates the

distance between each pair of atoms that have topologically equivalent atoms. These distances are the inter-atomic distance constraints for the topologically equivalent atoms in the model protein. In one embodiment, these distance constraints can be relaxed by accounting for the overall homology between the model and template protein as discussed below.

The protein modeler generates standard constraints based on the planarity of a side chain, the C-alpha and C-beta chirality, and C-alpha inter-atomic distances.

In a preferred embodiment, the protein modeler assigns a helical conformation to each amino acid in the model protein and establishes a position for each atom. The protein modeler then uses well-known software packages to generate chemical constraints such as inter-atomic distances between 1-2 and 1-3 neighbors based on the connectivity of the model protein. The inter-atomic distance constraints are used to constrain the topology relating to non-topologically equivalent atoms (non-TEAs). The selection of a helical conformation is arbitrary and used to establish position information for the standard software package. One skilled in the art would appreciate that one could generate such 1-2 and 1-3 neighbor constraints without use of standard software packages, and, thus obviate the need to establish the position of each atom to generate these constraints.

Once the various constraints are established, the well-known distance geometry software, DGEOM (discussed below), is invoked to calculate the position of each atom in the model protein and thus generates the three-dimensional structure of the model protein.

Figure 1 is an overall block diagram of the input and output for the protein modeler. The protein modeler 110 is preferably implemented on a computer system. In a preferred embodiment, the protein modeler 110 inputs template protein structure 111, the model protein partial structure 112, and the model/template protein sequence alignment. Based on the input information, the protein modeler generates a

model protein structure 114. The format of the information defining the protein structures are in a format defined by the Protein Data Bank (PDB). Table 1 illustrates the PDB protein structure format.

TABLE 1

Atom Portion							
	iAtom	Atom	AA	iAA	X	Y	Z
ATOM	1	N	ALA	1	0.000	0.000	0.000
ATOM	2	CA	ALA	1	0.762	-1.174	0.000
ATOM	3	C	ALA	1	0.638	-2.028	1.220
ATOM	4	O	ALA	1	0.589	-3.333	1.099
ATOM	5	CB	ALA	1	2.256	-0.891	-0.123
ATOM	6	N	ASP	2	0.569	-1.403	2.560
ATOM	7	CA	ASP	2	0.449	-2.161	3.767
ATOM	8	C	ASP	2	-0.904	-2.875	3.745
ATOM	9	O	ASP	2	-0.955	-4.068	4.177
ATOM	10	CB	ASP	2	0.524	1.274	4.963
ATOM	11	CG	ASP	2	1.899	-0.662	5.163
ATOM	12	OD1	ASP	2	2.906	-1.088	4.803
ATOM	13	OD2	ASP	2	1.801	0.439	5.828
ATOM	14	N	LEU	3	2.104	-2.178	3.231
ATOM	15	CA	LEU	3	-3.297	-2.981	3.276
ATOM	16	C	LEU	3	-3.207	-4.201	2.341
ATOM	17	O	LEU	3	-3.650	-5.307	2.689
ATOM	18	CB	LEU	3	-4.529	-2.195	2.856
ATOM	19	CG	LEU	3	-4.792	-0.924	3.666
ATOM	20	CD1	LEU	3	-6.093	-0.212	3.337
ATOM	21	CD2	LEU	3	-4.699	-1.311	5.152
ATOM	22	N	GLU	4	-2.587	-4.068	1.004
ATOM	23	CA	GLU	4	-2.545	-5.304	0.168
ATOM	24	C	GLU	4	-1.631	-6.304	0.886
ATOM	25	O	GLU	4	-1.925	-7.494	0.922
ATOM	26	CB	GLU	4	-1.990	-5.001	-1.237

Connectivity Portion					
		neighbor(1)	neighbor(2)	neighbor(3)	neighbor(4)
CONNECT	1	2	0	0	0
CONNECT	2	1	3	5	0
CONNECT	3	2	4	6	0
CONNECT	4	3	0	0	0
CONNECT	5	2	0	0	0
CONNECT	6	7	3	0	0
CONNECT	7	6	8	10	0
CONNECT	8	7	9	14	0
CONNECT	9	8	0	0	0
CONNECT	10	7	11	0	0
CONNECT	11	10	12	13	0
CONNECT	12	11	0	0	0
CONNECT	13	11	0	0	0
CONNECT	14	15	8	0	0
CONNECT	15	14	16	18	0
CONNECT	16	15	17	22	0
CONNECT	17	16	0	0	0

Table 1 contains an atom portion and a connectivity portion. The atom portion specifies for each atom in the protein a sequential atom number (iAtom), the atom name (Atom), the containing amino acid name (AA), the containing amino acid sequence number (iAA), and the Cartesian coordinates (position) of the atom (X,Y,Z). The connectivity portion indicates the connectivity of the atoms in the protein. The connectivity specifies each of the (up to four) neighbors of an atom.

Figure 2 is an overall flow diagram of the protein modeler. In step 201, the protein modeler generates an initial three-dimensional structure for the model protein. In a preferred embodiment, the protein modeler inputs model protein partial structure 112, which contains structure information in PDB format without position information. One skilled in the art would appreciate that the model protein partial structure 112 could be generated by the protein modeler based on the model protein amino acid sequence. The position information for each atom in the model protein is generated based on a helical conformation. This position information is used by DGEOM to determine chemical constraints between the atoms of the model protein. These chemical constraints are later used to constrain the position of non-topologically equivalent atoms. The arbitrary assignment of helical conformation to the peptide backbone and the choice of side chain rotomers does not impose any restriction on subsequent stages of the modeling procedure. In fact, non-helical conformations can also produce acceptable results. An acceptable helical conformation algorithm may be found Sundaram, K. and Srinivasan, S., Computer Programs Biomed, "Computer Simulated Modeling of Biomolecular Systems," Vol. 10, p. 29-34, 1979. As discussed above, this step is used to take advantage of predefined software packages that generate chemical constraints. However, the chemical constraints could be generated directly from the model protein amino acid sequence based on well-known chemical constraint principles such as 1-2 and 1-3 neighbor topology.

In step 202, the protein modeler generates a TEA position data structure for the model protein based on topologically equivalent atoms. The sequence alignment between the template and the model protein is used to generate the TEA position data structure for the protein model. The protein modeler retrieves the position information for each topologically equivalent atom in the template protein and stores the result in the TEA position data structure for the model protein. The protein modeler also indicates in the TEA position data structure which atoms have no topologically equivalent atom in the template protein. Table 2 illustrates a sample TEA position data structure (also known as a hybrid data structure because it contains information based on the model and template proteins). The TEA position data structure contains an entry for each atom in the model protein. Each entry contains the following fields. The derived field contains a 'T' for a topologically equivalent atom and an 'M' for a non-topologically equivalent atom. The fields model amino acid number (miAA), model amino acid name (mAA), model atom number (miAtom), and model atom name (mAtom) identify the atom of the model protein. The fields template amino acid number (tiAA), template amino acid name (tAA), template atom number (tiAtom), and template atom name (tAtom) identify the topologically equivalent atom in the template protein. The fields X, Y, and Z specify the position of a topologically equivalent atom.

TABLE 2

derived	m1AA	mAA	m1Atom	mAtom	t1AA	tAA	t1Atom	tAtom	X	Y	Z
T	1	ALA	1	N	9	LYS	65	N	2.905	16.102	16.260
T	1	ALA	2	CA	9	LYS	66	CA	3.912	15.058	16.725
T	1	ALA	3	C	9	LYS	67	C	5.266	15.385	16.010
T	1	ALA	4	O	9	LYS	68	O	6.290	15.212	16.692
T	1	ALA	5	CB	9	LYS	69	CB	3.466	13.692	16.221
T	2	ASP	6	N	10	LEU	74	N	5.295	15.639	14.727
T	2	ASP	7	CA	11	LEU	75	CA	6.466	16.097	13.966
T	2	ASP	8	C	10	LEU	76	C	7.251	17.154	14.667
T	2	ASP	9	O	10	LEU	77	O	8.444	17.104	15.130
T	2	ASP	10	CB	10	LEU	78	CB	6.037	16.475	12.514
T	2	ASP	11	CG	10	LEU	79	CG	7.066	16.512	11.446
T	2	ASP	12	OD1	10	LEU	80	OD1	6.325	16.645	10.067
T	2	ASP	13	OD2	10	LEU	81	OD2	7.874	17.787	11.570
T	3	LEU	14	N	11	ARG	82	N	6.616	18.280	14.957
T	3	LEU	15	CA	11	ARG	83	CA	7.108	19.506	15.459
T	3	LEU	16	C	11	ARG	84	C	7.571	19.194	16.081
T	3	LEU	17	O	11	ARG	85	O	8.732	19.516	17.254
T	3	LEU	18	CB	11	ARG	86	CB	6.052	20.688	15.582
T	3	LEU	19	CG	11	ARG	87	CG	6.569	21.939	16.339
T	3	LEU	20	CD1	11	ARG	88	CD	5.524	23.079	16.429
H	3	LEU	21	CD2							
T	4	GLU	22	N	12	GLN	93	N	6.717	18.621	17.675
T	4	GLU	23	CA	12	GLN	94	CA	7.107	18.413	19.062
T	4	GLU	24	C	12	GLN	95	C	8.174	17.335	19.110
T	4	GLU	25	O	12	GLN	96	O	8.879	17.329	20.161
T	4	GLU	26	CB	12	GLN	97	CB	5.948	18.162	20.019
T	4	GLU	27	CG	12	GLN	98	CG	5.015	19.373	20.013
T	4	GLU	28	CD	12	GLN	99	CD	3.833	19.168	21.006
T	4	GLU	29	OE1	12	GLN	100	OE1	3.439	18.011	21.258
T	4	GLU	30	OE2	12	GLN	101	NE2	3.240	20.354	21.278
T	5	ASP	31	N	13	GLY	102	N	8.109	16.289	18.305
T	5	ASP	32	CA	13	GLY	103	CA	9.140	15.202	18.416
T	5	ASP	33	C	13	GLY	104	C	10.525	15.821	17.951
T	5	ASP	34	O	13	GLY	105	O	11.502	15.199	18.495
T	5	ASP	35	CB							
T	5	ASP	36	CG							
T	5	ASP	37	OD1							
T	5	ASP	38	OD2							

In step 203, the protein modeler generates the distance constraints based on the TEA position data structure. The protein modeler determines the distance between each topologically equivalent atom. In step 204, the protein modeler generates standard constraints for non-topologically equivalent atoms. These standard constraints include planarity of side chains, 1-4 inter-C-alpha distances, and C-alpha and C-beta chirality. In step 205, the protein modeler

inputs miscellaneous constraints. These miscellaneous constraints allow for the methods of the present invention to be expanded to include other constraints as they become known. For example, a miscellaneous constraint could be distance constraints imposed by disulfide crosslinks. In step 206, the protein modeler generates the three-dimensional structure for the model protein based on the constraints and the helical three-dimensional structure of the model protein. The protein modeler preferably employs the well-known program DGEOM to generate the three-dimensional structure for the model protein. The DGEOM program is one of the various well-known programs that are based on the principles of distance geometry. Blaney et al., Quantum Chemistry Program Exchange, Program #590, (1989), Indiana University, Bloomington, Indiana, contains a description of DGEOM. A detailed explanation of the theory in practice of distance geometry can be found in Havel, T.F., Kuntz, I.D., and Crippen, G.M., Bull. Math Biol., "Theory and Practice of Distance Geometry," Vol. 45, pp. 666-720, 1983.

Figure 3 is a flow diagram of a routine to generate a TEA position data structure for the model protein. This routine inputs the model and template sequence alignment and the three-dimensional structure of template protein, and outputs the TEA position data structure for the model protein. In step 301, the routine selects the next model and template amino acid from the sequence alignment. In step 302, if all the amino acids have already been selected, then the routine returns, else the routine continues at step 303. In step 303, if the model protein has an amino acid in the sequence position but the template protein has no amino acid, then the amino acid is part of a variable region and the routine continues at step 304, else the routine continues at step 305. In step 304, the routine adds an entry to the TEA position data structure for each atom of the model amino acid as a non-topologically equivalent atom and loops to step 301 to select the next amino acid in the sequence alignment. In step 305, if both the model and template proteins have an amino acid in the sequence alignment, then the routine continues at step 306 to determine which atoms between the amino acids are

topologically equivalent, else the routine loops to step 301. In step 306, the routine selects the next atom in the model amino acid starting with a first atom. In step 307, if all the atoms in the model amino acid have already been selected, then the routine loops to step 301, else the routine continues at step 308. In step 308, if the template amino acid has a topologically equivalent atom to the selected atom of the model amino acid, then the routine continues at step 309, else the routine continues at step 310. In step 309, the routine adds an entry to the TEA position data structure for the selected atom as a topologically equivalent atom and loops to step 306. In step 310, the routine adds an entry to the TEA position data structure for the selected atom as a non-topologically equivalent atom and the routine loops to step 306.

CODE TABLE 1

```

void GenerateTEAPositionDataStructure()
{
    imAtom = 1;
    itAtom = 1;

    for (i=1; i<=cAminoAcids; i++)
        (tAA = Align(i).tAA;

        mAA = Align(i).mAA;

        if (tAA == '-' && mAA != "-")
            AddVarAA();

        if (tAA != "-" && mAA != "-")
            (for (j=1; j<=cmaxAtoms; j++)
                if (PIM[mAA][j] == 1
                    if (PIM[tAA][j] == 1)
                        AddTEA(imAtom++, itAtom++);
                    else
                        AddNTEA(imAtom++);
            )
        )
    )

void AddVarAA()
{
    do
        AddNTEA(imAtom++);
    while (Helical[imAtom].IAA == Helical[imAtom-1].IAA)
}

void AddNTEA(index x)
{
    Hybrid[x].derived = "M";
    Hybrid[x].imAA = Helical[x].IAA;
    Hybrid[x].mAA = Helical[x].AA;
    Hybrid[x].imAtom = Helical[x].iAtom;
    Hybrid[x].mAtom = Helical[x].Atom;
}

void AddTEA(index x, y)
{
    Hybrid[x].derived = "T";
    Hybrid[x].imAA = Helical[x].IAA;
    Hybrid[x].mAA = Helical[x].AA;
    Hybrid[x].imAtom = Helical[x].iAtom;
    Hybrid[x].mAtom = Helical[x].Atom;
    Hybrid[x].itAA = Templat[y].IAA;
    Hybrid[x].tAA = Templat[y].AA;
    Hybrid[x].itAtom = Templat[y].iAtom;
    Hybrid[x].tAtom = Templat[y].Atom;
    Hybrid[x].x = Templat[y].x;
    Hybrid[x].y = Templat[y].y;
    Hybrid[x].z = Templat[y].z;
}

```

Code Table 1 contains "C" programming language pseudocode corresponding to the flow diagram of Figure 3. The data structure Align contains the protein sequence alignment. The data structure Helical contains data relating to the three-dimensional structure of a helical conformation of the model protein. However, the pseudocode does not use the helical position data. The data structure Template contains the data relating to the three-dimensional structure of the template protein. The data structure Hybrid contains the TEA position data structure. The fields names are the same as described above for the three-dimensional structure in PDB format and the TEA positional data structure. The pseudocode also uses the position identity matrix (PIM) defined in Table 2.

Table 2 is the position identity matrix for side chain atoms of amino acids. The rows of table 2 represent amino acids and the columns represent positional identity of the amino acids. Topologically equivalent atoms in different amino acids are aligned in columns with an entry of 1. For example, if alanine is replaced by arginine in the model, the position of the arginine atoms--N, C-alpha, C', O, and C-beta--in the template protein are used. Although not shown, the position identity matrix also contains columns for each of the four backbone atoms.

TABLE 2

	B	G1	G2	D1	D2	E1	E2	E3	Z1	Z2	Z3	H1	H2
	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)	(15)	(16)	(17)
Ala	1	0	0	0	0	0	0	0	0	0	0	0	0
Arg	1	1	0	1	0	1	0	0	1	0	0	1	1
Asn	1	1	0	1	1	0	0	0	0	0	0	0	0
Asp	1	1	0	1	1	0	0	0	0	0	0	0	0
Cys	1	1	0	0	0	0	0	0	0	0	0	0	0
Glu	1	1	0	1	0	1	1	0	0	0	0	0	0
Gln	1	1	0	1	0	1	1	0	0	0	0	0	0
Gly	0	0	0	0	0	0	0	0	0	0	0	0	0
His	1	1	0	1	1	1	1	0	0	0	0	0	0
Ile	1	1	1	1	0	0	0	0	0	0	0	0	0
Leu	1	1	0	1	1	0	0	0	0	0	0	0	0
Lys	1	1	0	1	0	1	0	0	1	0	0	0	0
Met	1	1	0	1	0	1	0	0	0	0	0	0	0
Phe	1	1	0	1	1	1	1	0	0	0	0	0	0
Pro	1	1	0	1	0	0	0	0	0	0	0	0	0
Ser	1	1	0	0	0	0	0	0	0	0	0	0	0
Thr	1	1	1	0	0	0	0	0	1	0	0	0	0
Trp	1	1	0	1	1	1	1	1	0	1	1	1	0
Tyr	1	1	0	1	1	1	1	0	1	0	0	1	0
Val	1	1	1	0	0	0	0	0	0	0	0	0	0

Figure 4 is a flow diagram of the routine to generate distance constraints for topologically equivalent atoms. The routine inputs the TEA position data structure of the model protein and outputs distance constraints for the topologically equivalent atoms. In step 401, the routine selects the next topologically equivalent atom in the TEA position data structure, starting with the first topologically equivalent atom. In step 402, if all the topologically equivalent atoms have already been selected, then the routine returns, else the routine continues at step 403. In steps 403 through 407, the routine loops determining the distance between the selected topologically equivalent atom and all other topologically equivalent atoms. In step 403, the routine chooses a next topologically equivalent atom after the selected topologically equivalent atom in the TEA position data structure. In step 404, if all the topologically equivalent atoms after the selected topologically equivalent atom have already been chosen, then the routine loops to step 401 to select another topologically equivalent atom, else the

routine continues at step 405. In step 405, the routine calculates the distance between the chosen and selected topologically equivalent atoms. In step 406, the routine calculates the upper and lower bounds of the distance constraints. As discussed above, the upper and lower bounds can be relaxed based on the overall homology between the model and template proteins. In step 407, the routine outputs the distance constraints between the chosen and the selected topologically equivalent atoms and the routine loops to step 403. Code Table 2 contains "C" programming language pseudocode for the flow diagram of Figure 4.

CODE TABLE 2

```
void GenerateTEAConstraints()
{
    for (i=1; i<=nmAtoms; i++)
        if (Hybrid[i].derived == "T")
            for (j=i; j<=nmAtoms; j++)
                if (Hybrid[j].derived == "T")
                    (Calculate distance between atom i and j
                     OutputmAA and mAtom for i and j
                     upper and lower bounds of distance
                    );
}
```

Figure 5 is a flow diagram of a routine to generate standard constraints for non-topologically equivalent atoms. The routine inputs the TEA position data structure and outputs the standard constraints for non-topologically equivalent atoms. In step 501, the routine selects the next non-topologically equivalent atom in the TEA position data structure, starting with the first non-topologically equivalent atom. In step 502, if all the non-topologically equivalent atoms have already been selected, then the routine returns, else the routine continues at step 503. In step 503, if the selected non-topologically equivalent atom is part of a planar side chain, then the routine outputs a planar constraint in step 504 and continues at step 505. In step 505, if the selected non-topologically equivalent atom is a C-

beta atom and the amino acid is isoleucine or threonine, then the routine outputs a C-beta chiral constraint in step 506 and continues at step 507. In step 507, if the selected non-topologically equivalent atom is a C-alpha atom and the amino acid is not glycine, then the routine outputs a C-alpha chiral constraint in step 508 and continues at step 509. In step 509, if the selected non-topologically equivalent atom is a C-alpha atom, then the routine outputs the distance constraints for the next C-alpha atom in the peptide chain. The routine then loops to step 501 to select the next non-topologically equivalent atom. Code Table 3 contains "C" programming language pseudocode for the flow diagram of Figure 5.

CODE TABLE 3

```
void GenerateNTEAConstraints()
{
    for (i=1; i<nmAtoms; i++)
        if (Hybrid[i].derived == model)

            (if (Hybrid[i].MAA is planar)
                Output ("plane", ids of atoms in plane);

            if (Hybrid.mAtom is CB &&
                Hybrid[i].MAA is isoleucine or threonine)
                Output ("chiral", ids of side chain atoms,
                    chiral volume);

            if (Hybrid[i].mAtom is CA &&
                Hybrid[i].MAA is not glycine)
                Output ("chiral", id of CA atom,
                    ids of side chain atoms, chiral volume);

            if (Hybrid[i].mAtom is CA)
                Output (id of i atom, id of next CA atom,
                    upper and lower distance bound);

    }
}
```

Figure 6 is a flow diagram of the utilization of the DGEOM software package. In step 601, DGEOM initializes an inter-atomic distance matrix based on the helical three-dimensional structure of the model protein. DGEOM determines inter-atomic distances based on chemical constraints such as inter-atomic distances between 1-2 and 1-3 neighbors. Although DGEOM establishes these distance constraints for all atoms within the model protein, the distance constraints for topologically equivalent atoms are eventually overridden by distance constraints imposed by the template protein. In step 602, DGEOM applies the distance constraints for the topologically equivalent atoms to the distance matrix. In step 603, DGEOM applies the standard constraints for non-topologically equivalent atoms to the distance matrix. In step 604, the routine applies miscellaneous constraints to the distance matrix. In step 605, DGEOM smoothes the upper and lower bounds in the distance matrix. In step 606, DGEOM generates the final three-dimensional structure of the model protein based on the smoothed distance matrix.

The following example illustrates the methods of the present invention. Table 3 contains a sequence alignment for the murine interleukin-4 and human interleukin-4 proteins. In this example, the three-dimensional structure of the human interleukin-4 protein is known, and the three-dimensional structure of the murine interleukin-4 (the model protein) protein is to be modeled. In this example, a template protein is derived from the human interleukin-4 protein. The variable regions in the template protein are preferably expanded to allow for flexibility in modeling the transitions between conserved and variable regions.

Continuing with the example, the protein modeler sets the position of each atom in the model protein with a topologically equivalent atom in the template protein to the position of the atom in the template protein. For example, the first aligned amino acids for these model and template proteins are both leucine. Since each atom in the model amino acid has a topologically equivalent atom in the template amino

acid, the protein modeler sets the position of each atom. Once the position of each topologically equivalent atom in the model protein is established, then the protein modeler generates the inter-atomic distance constraints for each pair of atoms. The protein modeler then determines standard constraints for the non-topologically equivalent atoms. For example, since the first amino acid, histidine, in the model protein does not align with an amino acid in the template protein, each atom of histidine is a non-topologically equivalent atom. Thus, a planar constraint, a C-alpha chiral constraint, and an inter-C-alpha distance constraint are generated. The protein modeler then inputs miscellaneous constraints that are generated by a user of the protein modeler. These constraints may typically be based on disulfide crosslinks or other empirically determined constraints. The protein modeler also generates chemical constraints for the non-topologically equivalent atom. Finally, the protein modeler applies all the constraints using the principles of the distance geometry to arrive at a three-dimensional structure of the murine interleukin-4 protein.

TABLE 3

```

MURIL4: -----HGCDKNHLREIIIGILNEVTGEGTPCTEMDVNPVLTATKNT
HUMIL4: -EAEAHKCDIT-LQEIIKTLNSLTEQKTLCTELTVTDIFAASKDT
TEMPLT: -----LQEIIKTLNSLTEQKTLCTELTVTDIFAASKDT

MURIL4: TESELVCRASKVLRI FYLKHGK-TPCLKKNS-----SVLMELQ
HUMIL4: TEKETFCAATVLRQFYSHHEKDTRCLGATAQQFHRHKQLIRFLK
TEMPLT: TEKETFCAATVLRQFYSHHEK---CL-----LIRFLK

MURIL4: RLFRAFRC LDSSISCTM NESKSTSLKDFLESLSKSIQMMDYS----
HUMIL4: RLDRNLWGLAGLNSCPVKEADQSTLENFLERLKTIMREKYSKCSS
TEMPLT: RLDRNLWGLAGLNSCPVKEADQSTLENFLERLKTIMREKYSKCSS

```

In an alternate embodiment, the present invention provides a method and system for modeling a model protein that bears weak sequence identity to a template protein with a known three-dimensional structure. Proteins with weak sequence identity, but with similar three-dimensional

structure, are known as divergent proteins. In this embodiment, the template protein and the model protein are preferably divergent proteins. The positions of the backbone atoms of the structurally conserved regions of the model protein are allowed to deviate from the position of the topologically equivalent atoms in the template protein. The deviation, called a "divergence factor," is expressed by specifying that inter-atomic distances as ranges rather than a single value. These ranges allow the modeling to accommodate for differences between aligned amino acids that are not identical. That is, the position of the backbone atoms can be adjusted to accommodate the differences in the side chains.

In one embodiment, all inter-atomic distances between pairs of atoms are allowed to deviate within the divergence factor. In particular, the inter-atomic distances between atoms in the same structurally conserved region are allowed to deviate (intra-region deviation), and the inter-atomic distances between atoms in different structurally conserved regions are allowed to deviate (inter-region deviation). Alternatively, intra-region deviation is not allowed. By not allowing intra-region deviations, the backbone of each structurally conserved region is held rigid but is allowed to rotate relative to other structurally conserved regions.

In this alternate embodiment, the protein modeler generates a three-dimensional structure of the model protein based on a template protein with a known three-dimensional structure, an amino acid sequence alignment between the model protein and the template protein, and an identification of the structurally conserved regions. Since the sequence identity between the model protein and the template protein is assumed to be weak, accurate alignment and identification of structurally conserved regions can be achieved by the well-known techniques, such as, "secondary structure prediction" or "disulphide topology."

To model a divergent protein, the protein modeler performs several modeling iteration with various divergence

factors. After each iteration, the number of distance violations for that iteration is recorded. A distance violation preferably occurs when a distance constraint cannot be satisfied to within a margin of error, which is .5 Å. The margin of error can be increased or decreased to allow for flexibility in the modeling. In a preferred embodiment, the protein modeler starts with a divergence factor of 0 and increases the divergence factor by 1 Å for each iteration. As the number of iterations increases, the number of violations typically tends to level off. The model generated when the number of violations levels off is typically selected as the three-dimensional structure for the model protein.

Modeling Variable Regions

In another embodiment, the present invention provides a method and system for modeling the variable regions of a model protein using the ψ and ϕ angles between amino acid pairs of a template protein or a family of template proteins. A variable region modeler generates a collection of ψ and ϕ angle values for each amino acid pair in a family of proteins. The variable region modeler creates a matrix of amino acid pairs, such that, each entry in the matrix is a list of all ψ and ϕ angle values that are in the variable regions of the family. The variable region modeler models the variable region by randomly selecting a ψ and ϕ angle value from the matrix for each amino acid pair in the variable region and calculating the corresponding atomic position information (e.g. Cartesian coordinates).

Each variable region is bounded by a beginning structurally conserved region and an ending structurally conserved region. The variable region modeler generates the atomic position information for the amino acids in the beginning structurally conserved region from the ψ and ϕ angle values in the corresponding structurally conserved region of a template protein. The variable region modeler then generates atomic position information for the amino acids in the

variable region. The variable region modeler successively selects each pair of amino acids in the variable region. For each pair of amino acids, the variable region modeler selects, preferably randomly, an ψ and ϕ angle value from the corresponding list in the matrix. The variable region modeler then calculates the atomic positions of the amino acids using the selected ψ and ϕ angle values. After processing the variable region, the variable region modeler generates the atomic positions for the amino acids in the ending structurally conserved region using the ψ and ϕ angle values of the corresponding structurally conserved region in the template protein. Finally, the variable region modeler calculates the root mean square (rms) deviation between the ending structurally conserved region of the model protein and the template protein. The rms deviation indicates the degree to which the model for the variable region is consistent with the bounding structurally conserved regions.

In a preferred embodiment, the variable region modeler iteratively generates models and rms deviations for a variable region. The variable region modeler selects the variable region model with smallest rms deviation. As the number of iterations is increased, the likelihood of generating successively smaller rms deviations also increases. In the preferred embodiment, the selection of ψ and ϕ angle values is random, that is, a Monte Carlo method. Alternatively, other selection criterion can be used. For example, selection of the ψ and ϕ angle values for each pair on the amino acids that follow and proceed the pair in the model protein. Also, as discussed above, amino acid angle matrix is preferably generated from the variable regions in a family of proteins. Alternately, the matrix can be generated from proteins not in the same family. Also, a matrix can be generated for each corresponding variable region in a family of protein. Each matrix contains the ψ and ϕ angle values for the corresponding variable region in each protein in the family. One skilled in the art would appreciate that other techniques for generating the matrix may be used.

After the atomic positions for the variable regions in a model protein are generated, then the positions may be used by the protein modeler as constraints to improve the modeling process as described above.

Figure 7 is a diagram illustrating the ψ and ϕ angles between pairs of amino acids. The backbone atoms of an amino acid are planar, that is they are located within one plane. The ψ and ϕ angles measure the inter-amino acid rotation between the C-alpha and C atoms and the N and C-alpha atoms, respectively. These angles specify the three-dimensional structure of the protein.

Figure 8 is a diagram illustrating the amino acid angle matrix. The rows and columns of the matrix 801 correspond to each of the 20 standard amino acid. Each entry of the matrix is a list of ψ and ϕ angle values for the corresponding amino acid pairs. For example, the entry 802 contains the angle values for the amino acid pair asparagine followed by arginine.

Figure 9 is a flow diagram of a function that generates the amino acid angle matrix. The function inputs the three-dimensional structure of a template protein and updates the matrix based on the ψ and ϕ angles of the amino acids in the variable regions of the template protein. In step 901, the function identifies the variable regions in the template protein. All regions that are not secondary structural regions (which are identified in the PDB) are variable regions. In step 903 through 908, the function loops processing each variable region and within each variable region each pair of amino acids. In step 903, the function selects the next variable region in the template protein, starting with the first variable region. In step 904, if all the variable regions have already been selected, then the function is complete, else the function continues at step 905. In step 905, the function selects the next amino acid pair in the selected variable region, starting with the first. In step 906, if all the amino acid pairs in the selected variable region have already been selected, then the function loops to

step 903 to select the next variable region, else the function continues at step 907. In step 907, the function calculates the ψ and ϕ angles based on the position information of the selected amino acids. In step 908, the function adds the calculated ψ and ϕ angle values to the list of angles in the matrix at the entry corresponding to the pair of the selected amino acids. The function then loops to step 905 to select the next pair of amino acids.

Figure 10 is a flow diagram of a function to generate a variable region model for a model protein based on the amino acid angle matrix. The function outputs an rms deviation value for the generated model. In a preferred embodiment, this function is invoked multiple times, and the model with the smallest rms deviation is selected as the model for the variable region. In step 1001, the function sets the position information for the beginning structurally conserved region. The position information is generated from the ψ and ϕ angle values of the corresponding structurally conserved region in the template protein. Alternatively, the coordinates can be obtained directly from the three-dimensional structure of the template protein. In steps 1002 through 1005, the function loops selecting each pair of amino acids in the variable region, selecting ψ and ϕ angle values, and setting position information for the atoms of the selected amino acids. In step 1002, the function selects the next amino acid pair in the variable region, starting with the first. In step 1003, if all the amino acid pairs in the variable region have been selected, then the function continues at step 1006 to process the ending structurally conserved region, else the function continues at step 1004. In step 1004, the function selects an ψ and an ϕ angle value from the matrix at the entry corresponding to the selected amino acid pair. In a preferred embodiment, the ψ and ϕ angles are selected randomly from the list. Alternatively, the ψ and ϕ angles can be selected based on the probability of occurrence of the angles in the template protein or family of template proteins. Also, the ψ and ϕ angle values can be

selected based on a probability that certain ψ and ϕ angle values proceed or follow other ψ and ϕ angle values. In step 1005, the function calculates position information for each backbone atom in the selected pair of amino acids based on the angles and the calculated position information of the adjacent amino acid. The function also calculates the position information for the side chain atoms and loops to step 1002 to select the next pair of amino acids. In step 1006, the function determines the position information for the amino acids in the ending structurally conserved regions. The position information is calculated based on the calculated position of the last amino acid in the variable region and the ψ and ϕ angles for each pair of amino acids in the ending structurally conserved region of the template protein. In step 1007, the function calculates the rms deviation between the ending structurally conserved region of the model protein and the corresponding region in the template protein.

In alternate embodiment, the three-dimensional structure of a protein can be modeled de novo from the collected ψ and ϕ angle values. A matrix of ψ and ϕ angle values can be generated for all proteins in the PDB or for a family of proteins. The positional information can then be generated for each pair of amino acids in the model protein by selecting (e.g. randomly) ψ and ϕ angle values from the matrix. The compactness of the resulting three-dimensional structure can then be computed. The compactness can include a measurement of the volume of the three-dimensional structure, a measurement of the maximum inter-atomic distance, or a measurement of the density. The three-dimensional structure that is most compact can then be selected as the folded form of the model protein. Also, the ψ and ϕ angle values can be combined with other constraints as described above to model a protein.

Although the present invention has been described in terms of a preferred embodiment, it is not intended that the invention be limited to this embodiment. Modifications within

in the art, the scope of the present invention is defined by the claims which follow.

Claims

1. A method in a computer system for modeling a three-dimensional structure of a model protein, the modeling based upon a three-dimensional structure of a template protein and an amino acid sequence alignment of the model protein and the template protein, the proteins comprising a plurality of amino acids, each amino acid having backbone atoms and side chain atoms, each atom in a three-dimensional structure having a position, the method comprising the steps of:

for each amino acid in the model protein, when the template protein has an amino acid aligned with the amino acid of the model protein, establishing the position of each backbone atom of the amino acid of the model protein based on the position of a topologically equivalent backbone atom in the aligned amino acid of the template protein;

generating inter-atomic distance constraints for each pair of atoms with an established position; and

setting the position of each atom in the model protein wherein the inter-atomic distances are in accordance with the constraints.

2. The method of claim 1 including the step of generating standard constraints for amino acids of the model protein having an atom with no topological equivalent atom in the aligned amino acid of the template protein.

3. The method of claim 2 wherein the step of generating the standard constraints includes the step of generating a constraint specifying that a side chain is planar.

4. The method of claim 2 wherein the step of generating the standard constraints includes the step of generating a constraint specifying the chirality of a C-beta atom of an isoleucine or threonine amino acid.

5. The method of claim 2 wherein the step of generating the standard constraints includes the step of generating a constraint specifying the C-alpha chirality of an amino acid.

6. The method of claim 2 wherein the step of generating the standard constraints includes the step of generating a constraint specifying inter-atomic distances between 1-4 neighbor C-alpha atoms.

7. The method of claim 1 wherein the step of setting the position of each atom further includes the step of ensuring that the inter-atomic distance between a side chain atom and another atom in the same amino acid are in accordance with chemical constraints.

8. The method of claim 7 including the step of generating chemical constraints based on 1-2 neighbor inter-atomic distances.

9. The method of claim 7 including the step of generating chemical constraints based on 1-3 neighbor inter-atomic distances.

10. The method of claim 1 wherein the step of establishing the position further includes the step of determining whether an atom of the template protein is topologically equivalent to an atom of the model protein.

11. The method of claim 10 wherein the step of determining further includes the step of using a position identity matrix, the position identity matrix specifying topological equivalence of atoms in different amino acids.

12. The method of claim 1 wherein the step of generating the inter-atomic distance constraints further includes the step of allowing for deviation of topology

between the model protein and the template protein based on the overall homology of the proteins.

13. The method of claim 1 including the step of inputting miscellaneous constraints.

14. The method of claim 13 including the step of inputting constraints based on disulfide crosslinks.

15. The method of claim 1 including performing the steps for each chain of the model protein when the model protein has multiple peptide chains.

16. The method of claim 1 including the step of deriving the template protein from multiple proteins.

17. The method of claim 1 including the step of expanding variable regions of the template protein to allow for flexibility in the modeling of the transition from a conserved region to a variable region.

18. The method of claim 1 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of a plurality of side chain atoms of the amino acid of the model protein based on the position of topologically equivalent side chain atoms in the aligned amino acid of the template protein.

19. The method of claim 18 including the step of generating standard constraints for amino acids of the model protein having an atom with no topological equivalent atom in the aligned amino acid of the template protein.

20. The method of claim 19 wherein the step of generating the standard constraints includes the step of generating a constraint specifying that a side chain is planar.

21. The method of claim 18 wherein the step of establishing the position further includes the step of determining whether an atom of the template protein is topologically equivalent to an atom of the model protein.

22. The method of claim 18 wherein the step of generating the inter-atomic distance constraints further includes the step of allowing for deviation of topology between the model protein and the template protein based on the overall homology of the proteins.

23. The method of claim 18 including the step of inputting miscellaneous constraints.

24. The method of claim 18 including the step of inputting constraints based on disulfide crosslinks.

25. The method of claim 18 wherein the step of setting the position of each atom further includes the step of ensuring that the inter-atomic distance between a side chain atom and another atom in the same amino acid are in accordance with chemical constraints.

26. The method of claim 1 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of each of the side chain atoms of the amino acid of the model protein based on the position of a topologically equivalent side chain atom in the aligned amino acid of the template protein.

27. The method of claim 25 wherein the step of setting the position of each atom further includes the step of ensuring that the inter-atomic distance between a side chain atom and another atom in the same amino acid are in accordance with chemical constraints.

28. The method of claim 26 including the step of generating standard constraints for amino acids of the model protein having an atom with no topological equivalent atom in the aligned amino acid of the template protein.

29. The method of claim 28 wherein the step of generating the standard constraints includes the step of generating a constraint specifying that a side chain is planar.

30. The method of claim 26 wherein the steps of establishing the position further includes the step of determining whether an atom of the template protein is topologically equivalent to an atom of the model protein.

31. The method of claim 26 wherein the step of generating the inter-atomic distance constraints further includes the step of allowing for deviation of topology between the model protein and the template protein based on the overall homology of the proteins.

32. The method of claim 26 including the step of inputting miscellaneous constraints.

33. The method of claim 26 including the step of inputting constraints based on disulfide crosslinks.

34. A computer-implemented method for identifying a template protein for a family of topologically related proteins based on the modeling of a three-dimensional structure of a model protein, the proteins comprising a

plurality of amino acids, each amino acid having backbone atoms and side chain atoms, the template protein having a known three-dimensional structure, each atom having a position in a three dimensional structure, the method comprising the step of:

for each amino acid in a model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of each backbone atom of the amino acid of the model protein based on the position of topologically equivalent backbone atoms in the aligned amino acid of the template protein;

generating inter-atomic distance constraints for each pair of atoms with an established position;

setting the position of each atom in the model protein wherein the inter-atomic distances are in accordance with the constraints to generate a three-dimensional structure for the model protein; and

accessing the conformance of the generated three-dimensional structure with the rules of protein folding.

35. The method of claim 34 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of a plurality of side chain atoms of the amino acid of the model protein based on the position of topologically equivalent side chain atoms in the aligned amino acid of the template protein.

36. The method of claim 34 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of each of the side chain atoms of the amino acid of the model protein based on the position of a topologically equivalent side chain atom in the aligned amino acid of the template protein.

37. A computer-implemented method for obtaining a structurally accurate sequence alignment between a model protein and a template protein, the proteins comprising a plurality of amino acids, each amino acid having backbone atoms and side chain atoms, the template protein having a known three-dimensional structure, each atom having a position in a three dimensional structure, the method comprising the steps of:

for a plurality of sequence alignments of the model protein and the template protein performing the steps of:

for each amino acid in the model protein, when the template protein has an amino acid aligned with the amino acid of the model protein, establishing the position of each backbone atom of the amino acid of the model protein based on the position of topologically equivalent backbone atom in the aligned amino acid of the template protein;

generating inter-atomic distance constraints for each pair of atoms with an established position;

setting the position of each atom in the model protein wherein the inter-atomic distances are in accordance with the constraints to generate three-dimensional structure for the model protein; and

accessing the conformance of the generated three-dimensional structure with the rules of protein folding.

38. The method of claim 37 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of a plurality of side chain atoms of the amino acid of the model protein based on the position of topologically equivalent side chain atoms in the aligned amino acid of the template protein.

39. The method of claim 37 including the step of, for each amino acid in the model protein, when the template protein has an amino acid aligned with an amino acid of the model protein, establishing the position of each of the side

chain atoms of the amino acid of the model protein based on the position of a topologically equivalent side chain atom in the aligned amino acid of the template protein.

40. A method in a computer system for modeling a three-dimensional structure of a model protein, the model protein having amino acids, the amino acids having atoms, each atom having a position in the three-dimensional structure, the method comprising the steps of:

generating standard constraints for each amino acid in the model protein;

generating an inter-atomic distance matrix for the model protein in accordance with the constraints; and

setting the position of each atom in the model protein based on the distance matrix.

41. The method of claim 40 wherein the step of generating the standard constraints includes the step of generating a constraint specifying the C-alpha chirality of an amino acid.

42. The method of claim 40 wherein the step of generating the standard constraints includes the step of generating a constraint specifying inter-atomic distances between 1-4 neighbor C-alpha atoms.

43. The method of claim 40 wherein the step of setting the position of each atom further includes the step of ensuring that the inter-atomic distance between a side chain atom and another atom in the same amino acid are in accordance with chemical constraints.

44. A method in a computer system for modeling a three-dimensional structure of a model protein, the model protein having amino acids, the amino acids having atoms, each atom having a position in the three-dimensional structure, the method comprising the steps of:

inputting miscellaneous constraints for each amino acid in the model protein;

generating an inter-atomic distance matrix for the model protein in accordance with the constraints; and

setting the position of each atom in the model protein based on the distance matrix.

45. The method of claim 44 wherein the miscellaneous constraints are based on disulfide crosslinks.

46. A method in a computer system for modeling a three-dimensional structure of a variable region of a model protein, the model protein having amino acids, the amino acids having positions within a three-dimensional structure, the method comprising the steps of:

receiving relative positional information between pairs of amino acids;

establishing a position for a first amino acid of the variable region; and

for each amino acid pair in the variable region, generating a model position for the amino acids based on the received relative positional information for the pair of amino acids.

47. The method of claim 46 wherein the relative positional information includes ψ and ϕ angle values between pairs of amino acids.

48. The method of claim 47 including the step of, for each combination of pairs of amino acids in variable regions of a family of proteins, collecting the ψ and ϕ angle values for each pair of amino acids and wherein the step of generating a model position bases the model position on one of the collected ψ and ϕ angle values.

49. The method of claim 48 wherein the step of generating a model position bases the model position on a

randomly selected one of the collected ψ and ϕ angle values.

50. The method of claim 46 including the step of generating a model position for the amino acids of the adjacent structurally conserved regions based on relative position information between pairs of amino acids.

51. The method of claim 50 wherein the step of generating a model position for the amino acids of the adjacent structurally conserved region is based on the ψ and ϕ angle values in a corresponding region of a template protein.

52. The method of claim 50 including the step of comparing the generated model position for the amino acids of the adjacent structurally conserved region to positions in a corresponding region in a template protein to indicate effectiveness of the modeling.

53. A method in a computer system for modeling a three-dimensional structure of a variable region of a model protein, the model protein having amino acids, the method comprising the step of establishing positional information for the amino acids in the variable region based on ψ and ϕ angle values between pairs of amino acids in a template protein.

54. The method of claim 53 wherein the model protein has a structurally conserved region that is adjacent to the variable region, and including the steps of:

establishing positional information for the amino acids in the adjacent structurally conserved region based on the established positional information of the amino acids in the variable region and based on ψ and ϕ angle values in a corresponding structurally conserved region of the template protein; and

comparing the established positional information of the amino acids of the adjacent structurally conserved region to positional information of the corresponding structurally

conserved region of the template protein to measure the effectiveness of the modeling.

55. The method of claim 53 including the step of collecting ψ and ϕ angle values for pairs of amino acids in the template protein and wherein the step of establishing bases the positional information on the collected ψ and ϕ angle values.

56. The method of claim 55 wherein the step of establishing includes the step of randomly selecting a collected ψ and ϕ angle value.

57. The method of claim 55 wherein the step of collecting collects ψ and ϕ angle values for pairs of amino acids in a family of template proteins.

58. A method in a computer system for modeling a three-dimensional structure for a variable region of a model protein, the protein having amino acids, the variable region having a corresponding beginning structurally conserved region and a corresponding ending structurally conserved region, the method comprising the steps of:

- collecting ψ and ϕ angle values for pairs of amino acids in a family of template proteins;

- generating three-dimensional positional information for the amino acids in the beginning structurally conserved region;

- generating three-dimensional positional information for the amino acids in the variable region based on the collected ψ and ϕ angle values and based on the generated positional information for the beginning structurally conserved region;

- generating three-dimensional positional information for the amino acids in the ending structurally conserved region based on the generated positional information for the variable region and based on positional information for a

corresponding ending structurally conserved region in a template protein; and

comparing the generated positional information for the amino acids in the ending structurally conserved regions to positional information for the amino acids in the corresponding structurally conserved region in the template protein to indicate correctness of the model.

59. The method of claim 58 including the step of randomly selecting collected ψ and ϕ angle values when generating the positional information for the amino acids in the variable region.

60. The method of claim 58 including the step of repeating the steps of generating and comparing, and including the step of selecting, as the model of the variable region, generated positional information for the amino acids in the variable region when the generated positional information of the amino acids in the ending structurally conserved region most closely compares to positional information in the corresponding structurally conserved region in the template protein.

61. A method in a computer system for modeling a three-dimensional structure of a model protein, the model protein having amino acids, the method comprising the step of establishing positional information for the amino acids in the model protein based on ψ and ϕ angle values between pairs of amino acids in a template protein.

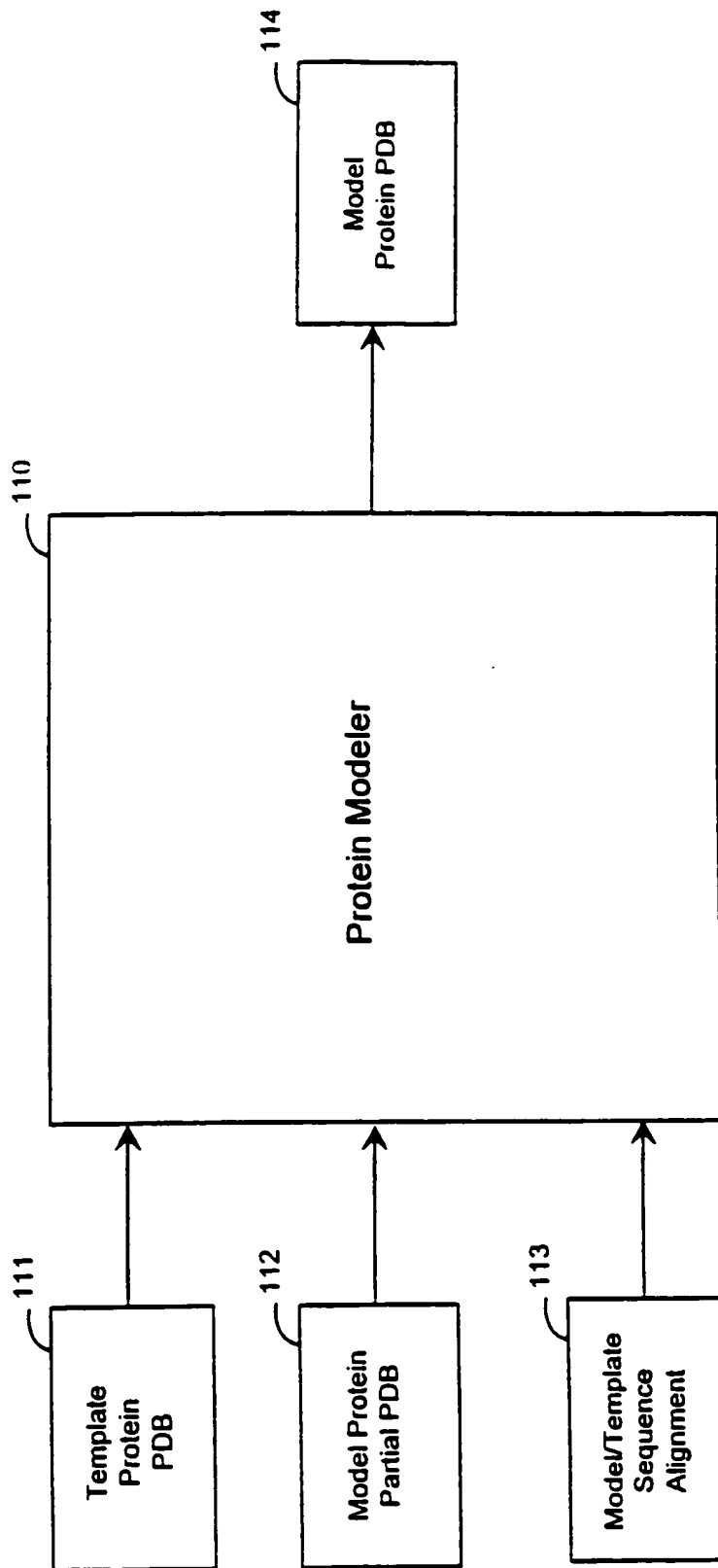
62. The method of claim 61 including the step of collecting ψ and ϕ angle values for pairs of amino acids in a template protein and wherein the step of establishing bases the positional information on the collected ψ and ϕ angle values.

63. The method of claim 1 wherein the step of

generating inter-atomic distance constraints for each pair of atoms includes the step of including a divergence factor into the distance constraints.

64. The method of claim 1 wherein the step of generating inter-atomic distance constraints for each pair of atoms includes the step of including a divergence factor into the distance constraints for inter-region distance and not including a divergence factor into the distance constraints for intra-region distances.

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**FIG. 1**

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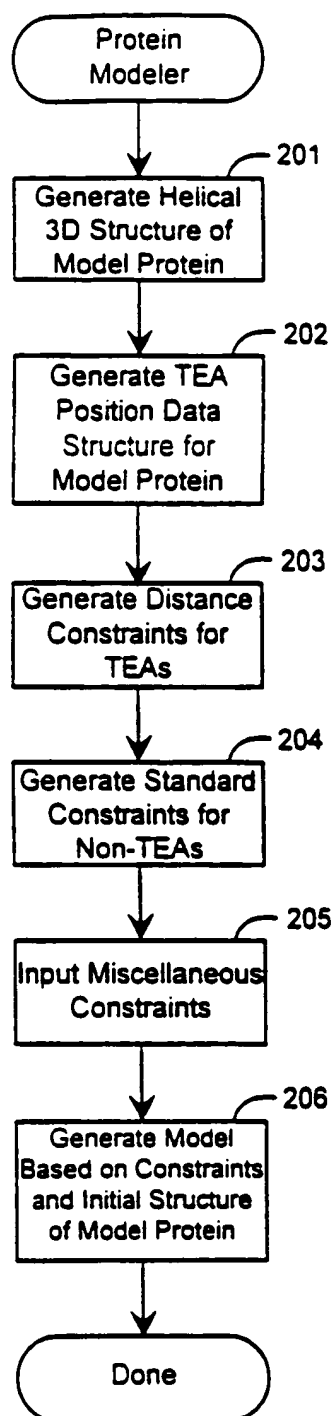
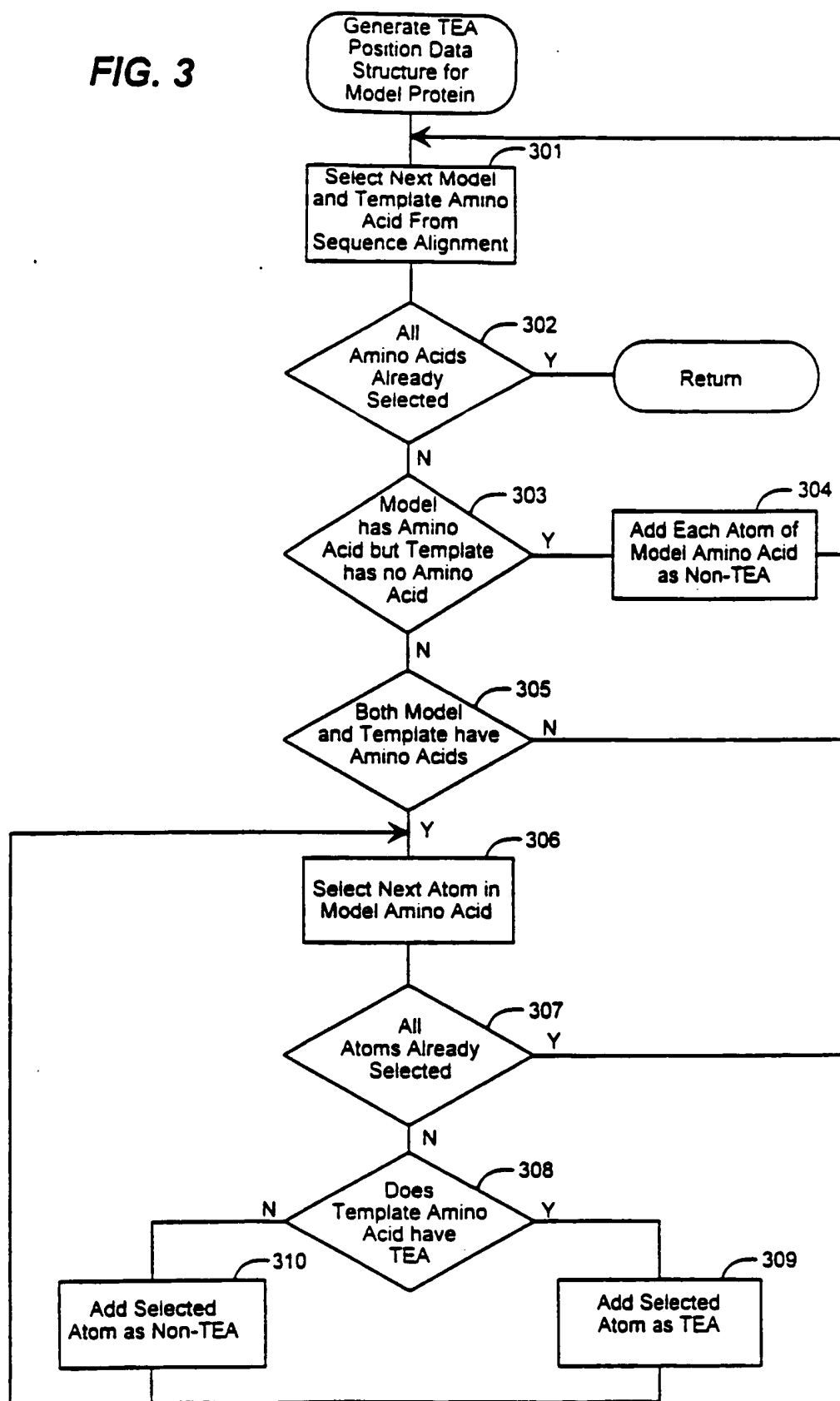
**FIG. 2**

FIG. 3



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FIG. 4

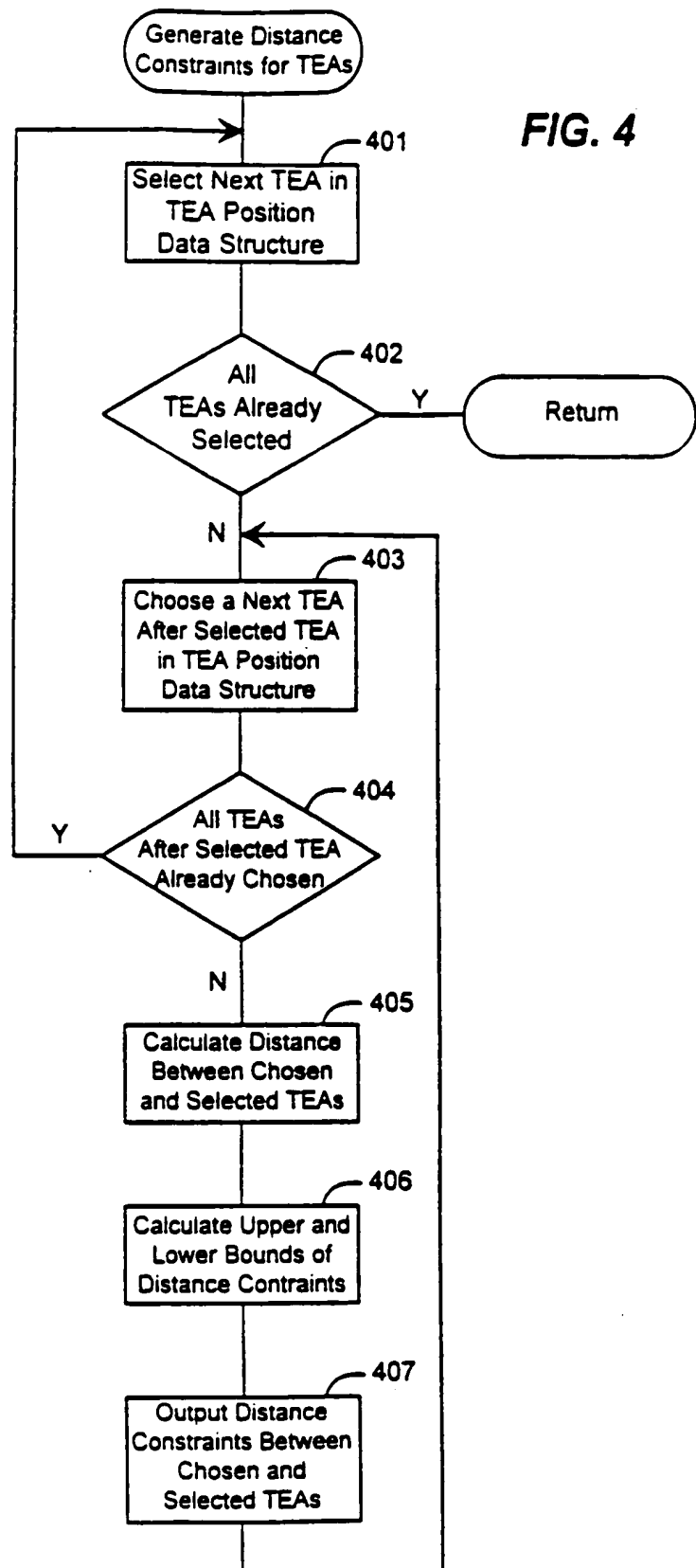
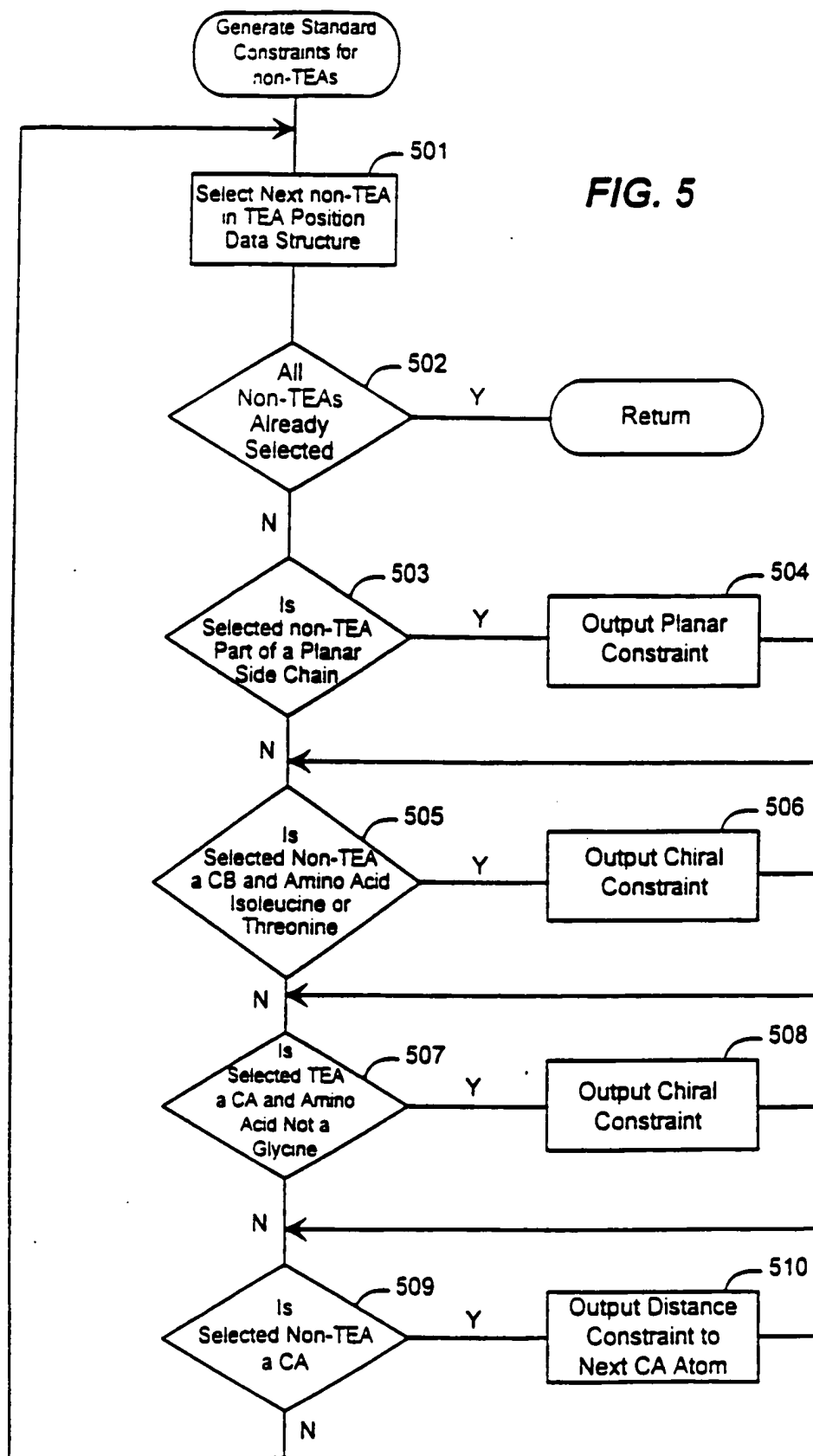
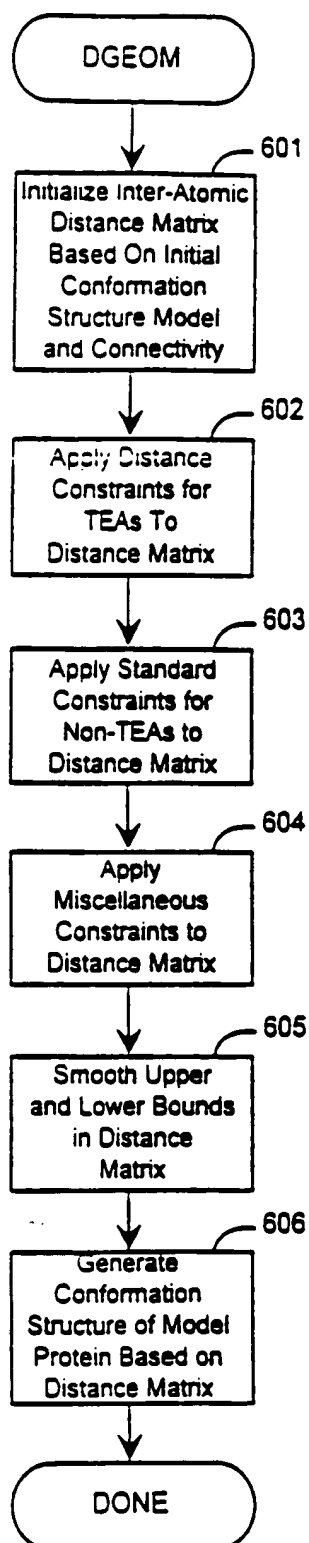


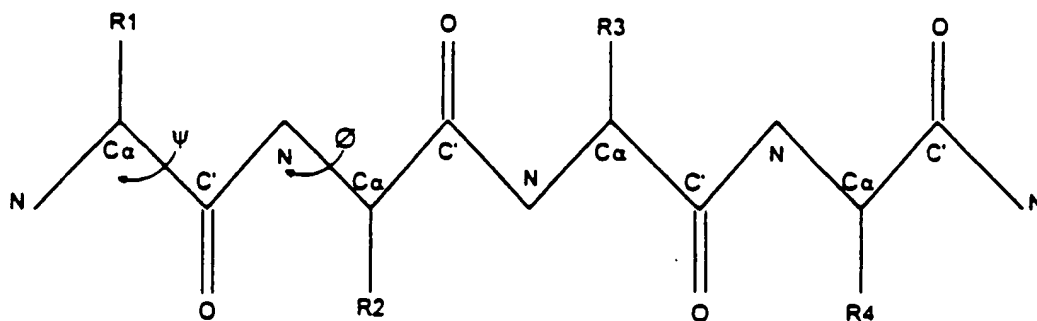
FIG. 5



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**FIG. 6**

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**FIG. 7****Amino Acid Angle Matrix**

	Ala	Arg	Asn	801
Ala				
Arg				
Asn				

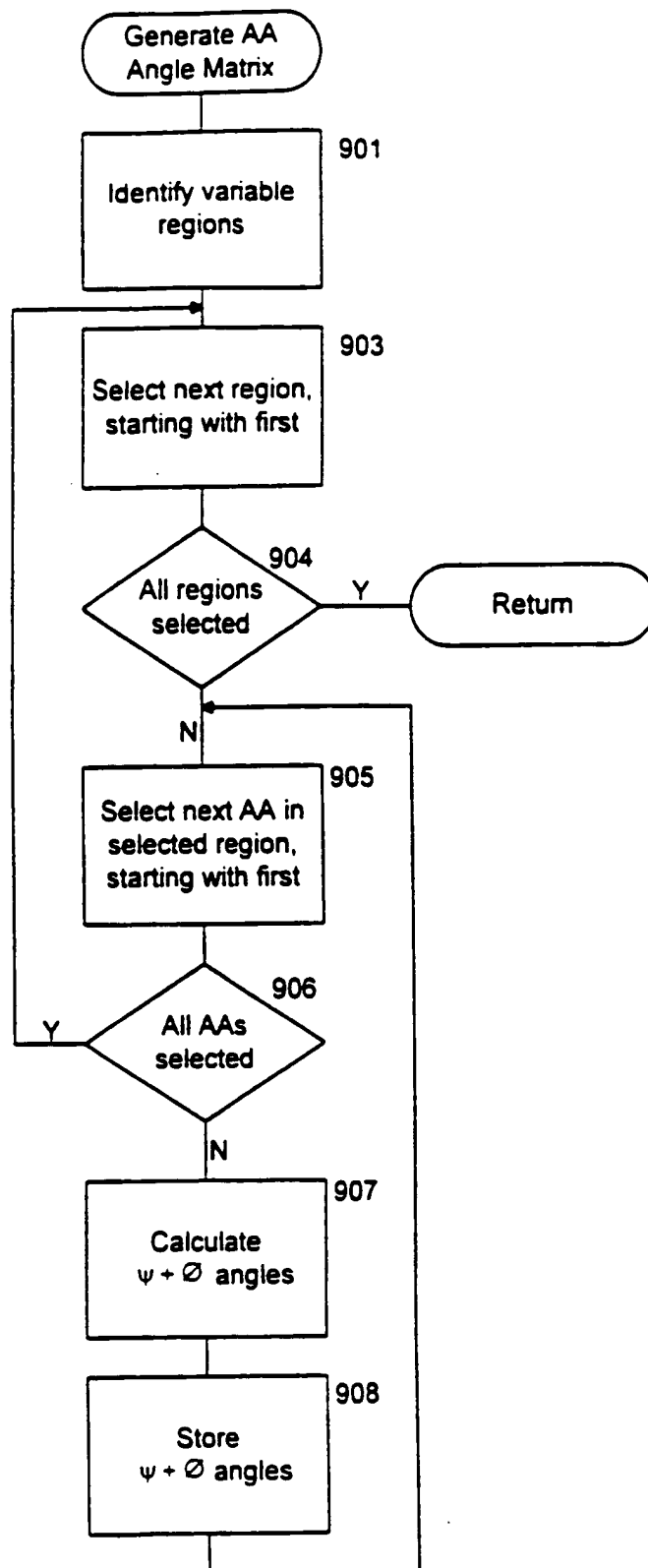
Angle List	
ψ	ϕ

802

FIG. 8

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**FIG. 9**

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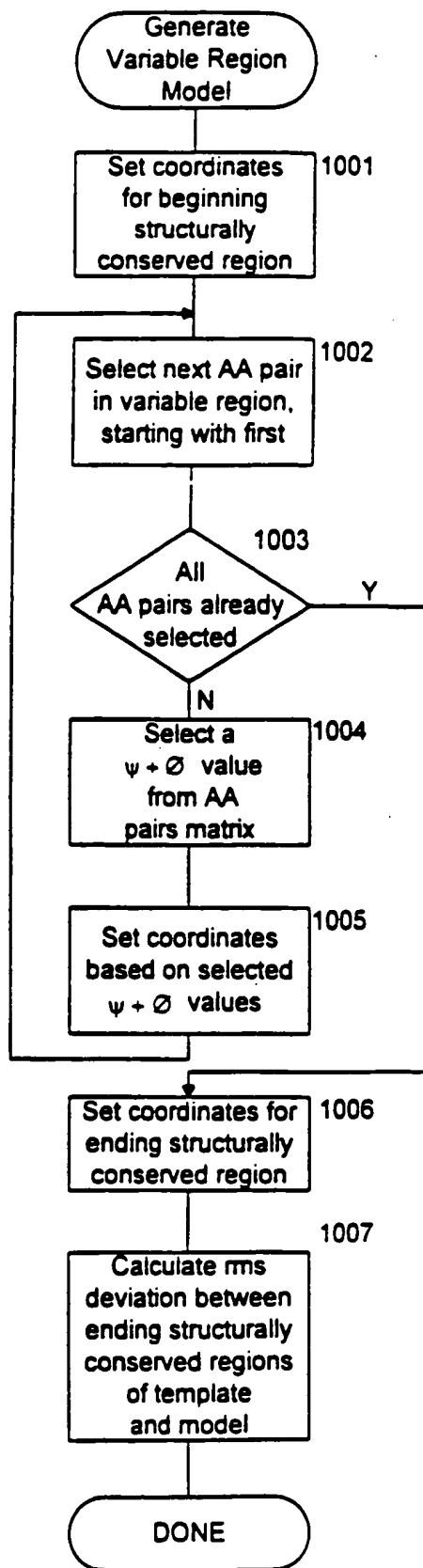


FIG. 10
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INTERNATIONAL SEARCH REPORT

International application No
PCT/US94/04822

A. CLASSIFICATION OF SUBJECT MATTER

IPC(5) : G01N 24/00

US CL : 364/496

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 364/496, 578

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y, P	US,A,5,241,470 (Lee et al.) 31 August 1993, col. 27, lines 58-60; col. 19, lines 46-50; col. 28, lines 36-38; col. 30, lines 53,54	1 - 5, 10, 13, 14, 19, 20, 21, 23, 24, 28, 29, 30, 33, 34, 36-41, 44, 45
Y	Journal of Computational Chemistry, Volume 4, No. 2, issued 1983, Brooks et al., "CHARMM: A Program for Macromolecular Energy, Minimization, and Dynamics Calculations, pages 187-217, especially page 196.	1-5, 10, 13, 14, 19, 20, 21, 23, 24, 28-30, 33, 34, 36-41, 44, 45

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 ☐ See patent family annex.

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O document referring to an oral disclosure, use, exhibition or other means		
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Date of the actual completion of the international search

07 JULY 1994

Date of mailing of the international search report

AUG 19 1994

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